Dated: February 26, 1997.

### Gene R. Haislip,

Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.

[FR Doc. 97–6919 Filed 3–19–97; 8:45 am] BILLING CODE 4410–09–M

### Manufacturer of Controlled Substances; Notice of Application

Pursuant to § 1301.43(a) of Title 21 of the Code of Federal Regulations (CFR), this is notice that on November 22, 1996, Knoll Pharmaceuticals, 30 North Jefferson Road, Whippany, New Jersey 07981, made application by renewal, which was received for processing February 4, 1997, to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of hydromorphone (9150), a basic class of controlled substance in Schedule II.

The firm plans to produce hydromorphone bulk product and finished dosage units of dilaudid for distribution to its customers.

Any other such applicant and any person who is presently registered with DEA to manufacture such substance may file comments or objections to the issuance of the above application.

Any such comments or objections may be addressed, in quintuplicate, to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, United States Department of Justice, Washington, D.C. 20537, Attention: DEA Federal Register Representative (CCR), and must be filed no later than May 19, 1997.

Dated February 26, 1997.

### Gene R. Haislip,

Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.

[FR Doc. 97–6917 Filed 3–18–97; 8:45 am] BILLING CODE 4410–09–M

# Manufacturer of Controlled Substances; Notice of Application

Pursuant to § 1301.43(a) of Title 21 of the Code of Federal Regulations (CFR), this is notice that on January 27, 1997, Mallinckrodt Chemical, Inc., Mallinckrodt & Second Streets, St. Louis, Missouri 63147, made application by renewal to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of the basic classes of controlled substances listed below:

Drug	Schedule
Tetrahydrocannabinols (7370)	1

Drug	Schedu
Methylphenidate (1724)	
Fentanyl (9801)	II

The firm plans to manufacture the controlled substances for distribution as bulk products to its customers.

Any other such applicant and any person who is presently registered with DEA to manufacture such substances may file comments or objections to the issuance of the above application.

Any such comments or objections may be addressed, in quintuplicate, to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, United States Department of Justice, Washington, DC 20537, Attention: DEA Federal Register Representative (CCR), and must be filed no later than May 19, 1997

Dated: February 26, 1997.

#### Gene R. Haislip,

Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.

[FR Doc. 97–6918 Filed 3–18–97; 8:45 am]

## Office of Justice Programs

### **Bureau of Justice Statistics**

Agency Information Collection Activities: Extension of a Currently Approved Collection; Comments Requested

**ACTION:** Notice of information collection under review; National corrections reporting program.

The Department of Justice, Office of Justice Programs, Bureau of Justice Statistics, has sent the following information collection to the Office of Management and Budget (OMB) for review and clearance in accordance with the review procedures of the Paperwork Reduction Act of 1996. The proposed information collection (60 day notice) was published in the **Federal Register** on January 13, 1997, to obtain comments from the public and affected agencies.

The purpose of this notice is to publish the collection for an additional 30 days until April 18, 1997. Comments should be directed to OMB, Office of Information and Regulatory Affairs, Attention: Department of Justice Desk Officer, Washington, DC 20503.

Request written comments and suggestions from the public and affected agencies concerning the proposed collection of information. Your comments should address one or more of the following four points:

(1) Evaluate whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information will have practical utility;

(2) Evaluate the accuracy of the agencies estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used;

(3) Enhance the quality, utility, and clarity of the information to be collected; and

(4) Minimize the burden of the collection of information on those who are to respond, including through the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology, e.g., permitting electronic submission of responses.

Overview of this information collection:

- (1) Type of Information Collection: Extension of a currently approved collection.
- (2) Title of the Form/Collection:
  National Corrections Reporting Program.
- (3) Agency form number, if any, and the applicable component of the Department of Justice sponsoring the collection: Form: NCRP-1A Prison Admissions Report; NCRP-1B Prison Release Report; NCRP-1C Parol Exit Report. Bureau of Justice Statistics, Office of Justice Programs, United States Department of Justice.
- (4) Affected public who will be asked or required to respond, as well as a brief abstract: Primary: State juvenile corrections agencies. Other: None. This